10/795,863

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L1 STR

Structure attributes must be viewed using STN Express query preparation.

L3 6 SEA FILE=REGISTRY SSS FUL L1

L4 1 SEA FILE=CAPLUS L3

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L4 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:756707 CAPLUS

DOCUMENT NUMBER: 141:277497

TITLE: Preparation of benzoylureidopyridylpiperidines for the

treatment of type 2 diabetes

INVENTOR(S): Schoenafinger, Karl; Kadereit, Dieter; Defossa,

Elisabeth; Herling, Andreas; Klabunde, Thomas

PATENT ASSIGNEE(S): Aventis Pharma Deutschland G.m.b.H., Germany

SOURCE: PCT Int. Appl., 33 pp.

CODEN. DIVVDO

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

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WO 2004078743
                          A1
                                 20040916
                                             WO 2004-EP1735
                                                                     20040221
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             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
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PRIORITY APPLN. INFO.:
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                                             US 2003-487497P
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                                                                     20030715
                                             WO 2004-EP1735
                                                                  A 20040221
OTHER SOURCE(S):
                         MARPAT 141:277497
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AB Title compds. I [R1, R2 = H, halo, alkyl, etc.; R3 = H, alkyl, O-alkyl, etc.; X = OH, O-alkyl, NH2, etc.; A, B, D, E = CH, N, with the proviso that one of A, B, D or E is N; Y = (CH2)m; m = 0-2] and their pharmaceutically acceptable salts were prepared For example, condensation of amine II, e.g., prepared from 2-chloro-3-nitropyridine in 2-steps, and

III

2-chloro-4-fluorobenzoylisocyanate, afforded ureidopyridylpiperidine III. In activated glycogen phosphorylase inhibition assays, 4-examples of compds. I exhibited IC50 values ranging from 0.01-3.65 \(\mu M \), the IC50 value of benzoylurea III was 0.04 μM. Compds. I were claimed useful for the treatment of type 2 diabetes. 758720-48-8P 758720-49-9P 758720-50-2P 758720-51-3P 758720-52-4P 758720-53-5P RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses) (preparation of benzoylureidopyridylpiperidines for the treatment of type 2 diabetes) RN758720-48-8 CAPLUS CN 4-Piperidinecarboxylic acid, 1-[3-[[[(2-chloro-4fluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]- (9CI) (CA INDEX NAME)

RN 758720-51-3 CAPLUS

CN 4-Piperidinecarboxamide, 1-[3-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]- (9CI) (CA INDEX NAME)

RN 758720-52-4 CAPLUS

CN 4-Piperidinecarboxamide, 1-[3-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]-N-(2-hydroxyethyl)-(9CI) (CA INDEX NAME)

$$\begin{array}{c} & & & \\ & &$$

RN 758720-53-5 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[3-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]-4-phenyl- (9CI) (CA INDEX NAME)

10/795,863

REFERENCE COUNT:

THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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FILE 'USPAT2' ENTERED AT 14:55:07 ON 31 JAN 2007
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L1 STR

Structure attributes must be viewed using STN Express query preparation.

L3 6 SEA FILE=REGISTRY SSS FUL L1

L5 1 SEA L3

=> d 15 ibib abs hitstr

L5 ANSWER 1 OF 1 USPATFULL on STN

ACCESSION NUMBER: 2004:335674 USPATFULL

TITLE: Substituted benzoylureidopyridylpiperidine-and-

pyrrolidinecarboxylic acid derivatives, processes for

preparing them and their use

INVENTOR(S): Schoenafinger, Karl, Alzenau, GERMANY, FEDERAL REPUBLIC

OF

Kadereit, Dieter, Kelkheim, GERMANY, FEDERAL REPUBLIC

OF

Defossa, Elisabeth, Idstein, GERMANY, FEDERAL REPUBLIC

OF

Herling, Andreas, Bad Camberg, GERMANY, FEDERAL

REPUBLIC OF

Klabunde, Thomas, Frankfurt, GERMANY, FEDERAL REPUBLIC

OF

PATENT ASSIGNEE(S): Aventis Pharma Deutschland GmbH, Frankfurt am Main,

GERMANY, FEDERAL REPUBLIC OF (non-U.S. corporation)

NUMBER KIND DATE

US 2004266768 A1 20041230

PATENT INFORMATION: US 2004266768 A1 20041230 APPLICATION INFO.: US 2004-795863 A1 20040308 (10)

NUMBER DATE

PRIORITY INFORMATION: DE 2003-10309929 20030307

US 2003-487497P 20030715 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: ROSS J. OEHLER, AVENTIS PHARMACEUTICALS INC., ROUTE

202-206, MAIL CODE: D303A, BRIDGEWATER, NJ, 08807

NUMBER OF CLAIMS: 14 EXEMPLARY CLAIM: 1

LINE COUNT: 703

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention relates to compounds of the formula I ##STR1##

where the radicals are as defined, and their physiologically tolerated salts. The compounds are suitable, for example, as medicaments for preventing and treating type 2 diabetes.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 758720-48-8P 758720-49-9P 758720-50-2P

758720-51-3P 758720-52-4P 758720-53-5P

(preparation of benzoylureidopyridylpiperidines for the treatment of type 2 diabetes)

RN 758720-48-8 USPATFULL

CN 4-Piperidinecarboxylic acid, 1-[3-[[[(2-chloro-4-

fluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]- (9CI) (CA INDEX NAME)

RN 758720-49-9 USPATFULL

CN 4-Piperidinecarboxylic acid, 1-[3-[[[(2-chloro-4,5difluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]- (9CI) (CA INDEX NAME)

RN 758720-50-2 USPATFULL

CN 4-Piperidinecarboxylic acid, 1-[3-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]-, methyl ester (9CI) (CA INDEX NAME)

RN 758720-51-3 USPATFULL

CN 4-Piperidinecarboxamide, 1-[3-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]- (9CI) (CA INDEX NAME)

RN 758720-52-4 USPATFULL

CN 4-Piperidinecarboxamide, 1-[3-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]-N-(2-hydroxyethyl)-(9CI) (CA INDEX NAME)

RN 758720-53-5 USPATFULL
CN 4-Piperidinecarboxylic acid, 1-[3-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-2-pyridinyl]-4-phenyl- (9CI) (CA INDEX NAME)

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